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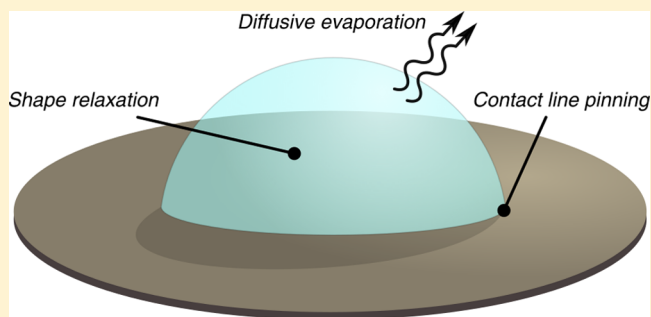
Macroscopic Model for Sessile Droplet Evaporation on a Flat Surface

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ABSTRACT: Evaporation of sessile droplets on a flat surface involves a complex interplay between phase change, diffusion, advection, and surface forces. In an attempt to significantly reduce the complexity of the problem and to make it manageable, we propose a simple model hinged on a surface free-energy-based relaxation dynamics of the droplet shape, a diffusive evaporation model, and a contact line pinning mechanism governed by a yield stress. Our model reproduces the known dynamics of droplet shape relaxation and of droplet evaporation, both in the absence and in the presence of contact line pinning. We show that shape relaxation during evaporation significantly affects the lifetime of a drop. We find that the dependence of the evaporation time on the initial contact angle is a function of the competition between the shape relaxation and evaporation and is strongly affected by any contact line pinning.



INTRODUCTION

Understanding the dynamics of spreading and drying of droplets deposited on a substrate is of importance to many practices, such as inkjet printing,^{1–3} pesticide spraying,⁴ and semiconductor device manufacturing.^{5,6} In the semiconductor industry, photolithographic methods are employed to define patterns for integrated circuits on wafers, coated with photosensitive polymer layers.^{5,7} Often, water immersion is used to increase the resolution of the lithography process.^{7,8} However, if any droplets are left behind on a wafer, they may induce the so-called watermark defects in the photoresist layer.^{5,9}

Due to the importance of understanding the drying processes, the drying of droplets on surfaces has been intensely studied experimentally,^{6,10–15} theoretically,^{15–19} and numerically.^{20–23} Nevertheless, the understanding of this multifaceted problem remains incomplete due to the multitude of coupled processes that determine the evaporation dynamics. Apart from the evaporation itself, processes such as convection and heat transport in the droplet, shape relaxation, and contact line pinning play a role.

Associated with the complex physics of the problem at hand are a large number of physical parameters, the relative importance of which depends on the initial and boundary conditions as well as the time and length scales of interest. Therefore, we aim to develop a macroscopic model that does not resolve the details of, for example, the velocity field inside the droplet or the vapor concentration field around it. Rather, we consider three constituents to make up our model: (1) interfacial free-energy-based relaxation for the droplet shape, (2) diffusion-limited evaporation, and (3) contact line pinning.

In the literature, various authors studied the evaporation of droplets, focusing on two limiting modes of evaporation: a

droplet evaporates with either a constant contact area or a constant contact angle, allowing transitions between these limits.^{13,16,24,25} Others have investigated the shape relaxation of droplets by measuring the contact angle of nonevaporating droplets in time.^{26–28} To combine the both aspects, which are described in the literature separately, we propose a model that is not restricted to the two evaporation modes, but contains the shape relaxation of the droplet during the evaporation process. It captures and extends the evaluation of Stauber et al.²⁴ by taking into account the contact line dynamics, i.e., incorporating both advancing and receding contact lines and considering cases without contact line pinning. Moreover, to describe the transition between mobile and pinned contact lines, our model includes a yield stress that governs contact line pinning: contact line motion is inhibited for capillary driving forces below a critical stress.

The remainder of this paper is organized as follows. In the **Theory** section, we present the main ingredients of our phenomenological model. The **Results and Discussion** section compares experimental data and existing theories, as well as presents an overview of representative cases of evaporation with and without contact line pinning. We also discuss in detail the implications of choices made for certain parameters during the calculations. In the **Summary and Conclusions** section, we summarize our results and present our main conclusions.

THEORY

The focus of this work is on droplets of sizes smaller than the capillary length $l_c = \sqrt{\gamma_{LG}/\rho g}$, which allows us to model the

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droplet as a spherical cap.²⁹ Here, γ_{LG} denotes the surface tension of the liquid–gas interface, ρ is the mass density of the fluid, and g is the gravitational acceleration. For water in air at room temperature, $l_c \simeq 3$ mm.²⁹ We presume the liquid to be incompressible. If the shape of the droplet is described as a spherical cap, it is uniquely defined by only two parameters. We choose for these the radius a of the contact area and the contact angle θ of the drop with the solid surface that we

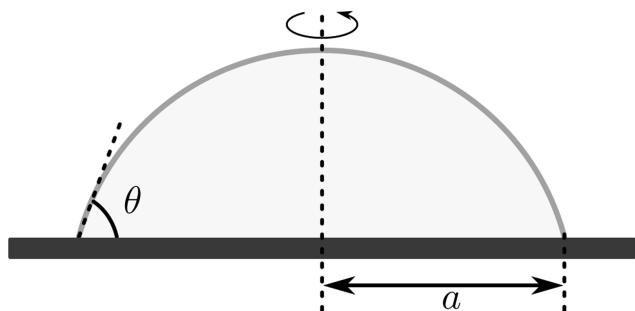


Figure 1. Schematic of an axisymmetric droplet on a planar surface, with contact angle θ and radius a of the contact area.

assume to be rigid (see Figure 1).³⁰ Geometrically, they are related to the droplet volume V according to

$$V(a, \theta) = \pi a^3 \left(\frac{2 - 3 \cos \theta + \cos^3 \theta}{3 \sin^3 \theta} \right) \quad (1)$$

Equation 1 implies that, for a given volume V , a prescribed value for the contact area radius a defines the contact angle θ and vice versa. Within a macroscopic description of the droplet shape, the equilibrium values of a and θ are determined by γ_{LG} , as well as by the solid–liquid and solid–gas interfacial tensions, γ_{SL} and γ_{SG} , respectively. We associate the droplet shape with an interfacial free energy $F(a, \theta)$, given by the sum of the interfacial tensions multiplied by the respective surface areas

$$F(a, \theta) = \pi a^2 \left(\gamma_{SL} - \gamma_{SG} + \frac{2\gamma_{LG}}{1 + \cos \theta} \right) \quad (2)$$

For a given volume V , eq 2 can be expressed as a function of θ only, using eq 1, that is, $F(a, \theta) \rightarrow F(\theta)$.³¹ Minimizing this free energy F with respect to θ produces the following equation for its optimal value

$$\gamma_{SG} - \gamma_{SL} - \gamma_{LG} \cos \theta_{eq} = 0 \quad (3)$$

which is the well-known Young’s equation for the equilibrium contact angle θ_{eq} .^{32,33} We note that θ_{eq} denotes the contact angle corresponding to the free-energy minimum; however, this angle is not necessarily straightforwardly accessible experimentally.³⁴

Out of equilibrium, eq 3 does not hold. To describe how an out-of-equilibrium droplet shape relaxes toward equilibrium, we construct a kinetic equation for the contact angle using a relaxational dynamics approach based on our free-energy landscape.^{35,36} It describes how the droplet adjusts its contact angle θ with the surface to move toward equilibrium. Together with the volume V , this defines a new radius of the contact area a . We refer to this process as “shape relaxation”, since both θ and a change simultaneously to accommodate a lower free-energy state. We note, however, that the overall shape of the

droplet remains a spherical cap. This allows us to quite naturally include the effects of steady evaporation and of a potential pinning of the contact line. In the next subsections, we discuss separately and in detail, the three main components of our phenomenological model: the relaxation dynamics of the droplet shape, the description for diffusive evaporation, and the contact line pinning mechanism.

Relaxation Dynamics of the Droplet Shape. On the basis of the free energy $F(\theta)$ obtained from eqs 1 and 2, we construct a relaxation equation for the contact angle θ

$$\frac{d\theta}{dt} = -\Gamma \frac{dF}{d\theta} \quad (4)$$

where Γ is a phenomenological relaxation rate that we specify in more detail below. Equation 4 describes a rate of change that is proportional to the generalized force $dF/d\theta$. This is in analogy to the so-called model A dynamics commonly applied in the kinetics of phase transitions of nonconserved order parameters.^{35–37} We note that it is also possible to derive kinetic equations by equating capillary and viscous forces.^{38,39} In our model, this balance is implicit in the parameter Γ .

Several experimental and theoretical works have identified the difference between the cosines of the instantaneous, time-dependent contact angle θ and its equilibrium value θ_{eq} , given by eq 3, to be the driving force for the motion of the contact line.^{27,28,33,40,41} For small values of the difference $\cos \theta - \cos \theta_{eq}$, this relaxation can be described by a simple exponential function.^{27,42} The exponential decay allows us to identify a characteristic time scale τ_{rx} as

$$\cos \theta - \cos \theta_{eq} \propto \exp(-t/\tau_{rx}) \quad (5)$$

We discuss the functional expression for τ_{rx} below.

To relate to eq 5, we transform eq 4 into a kinetic equation for $\cos \theta$ and expand it around the equilibrium $\cos \theta_{eq}$. A linearization produces an exponentially decaying $\cos \theta$, from which we determine Γ . This yields

$$\Gamma = \frac{1}{\tau_{rx} \gamma_{LG} V^{2/3} \Theta(\cos \theta_{eq}) \alpha(t)} \quad (6)$$

with

$$\Theta(x) \equiv 2(9\pi)^{1/3} (1-x)^3 (1+x) (2-3x+x^3)^{-5/3} \quad (7)$$

Equations 4 and 6 reproduce eq 5 for small deviations from equilibrium. In eq 6, we introduce a time-dependent, dimensionless factor $\alpha(t)$ to account for changes in the dynamics of the droplet shape relaxation due to, e.g., a change in droplet size over time. We return to this below. Equations 4 and 6 describe how a droplet deposited with a nonequilibrium initial contact angle θ_0 relaxes to the equilibrium value θ_{eq} in a relaxation process characterized by a fundamental time scale τ_{rx} , provided that the contact line is not pinned. We discuss our model for contact line pinning below.

The characteristic time scale τ_{rx} for the shape relaxation has been identified in various experimental and theoretical works to be dependent on the fluid viscosity η , the liquid–gas interfacial tension γ_{LG} , and a length scale L .^{19,26,27,39,41–44} as

$$\tau_{rx} = \frac{\eta L}{\gamma_{LG}} \quad (8)$$

In experiments on spreading of polymeric fluids, this length scale L has been described as a measure of the slip or friction length of the interaction between a polymeric liquid and the

solid,^{26,27,42} which seems to be independent of droplet dimensions²⁷ and has been estimated to be of the order of micrometers.⁴² In works on the coalescence of droplets, however, the length scale L has been shown to be proportional to the droplet size R ,^{19,39,41} which seems in agreement with the experimental and theoretical works on the spreading of polymer melts⁴³ and spherical droplets of simple liquids.⁴⁴ For this reason, we take the length scale L to be $kV_0^{1/3}$, with V_0 the initial volume of the droplet, making it proportional to the droplet size, and k a dimensionless proportionality constant that can be related to an Arrhenius factor.⁴¹ Hence

$$\tau_{\text{rlx}} = k \frac{\eta V_0^{1/3}}{\gamma_{\text{LG}}} \quad (9)$$

As the droplet size decreases during evaporation, the length scale L related to the shape relaxation may (1) remain constant (in the case that L is related to a slip or friction length) or (2) decrease with the droplet size. The scale factor $\alpha(t)$ can be employed to incorporate either behavior into the dynamics described by eqs 4 and 6. If L remains constant, we may choose $\alpha = 1$, whereas for a size-dependent length scale, $\alpha(t) = (V(t)/V_0)^{1/3}$. As we shall see, it turns out that the two expressions for α give rise to small differences in the droplet dynamics albeit that the lifetime of an evaporating droplet is not sensitive to whether α is proportional to the droplet size or not. For simplicity, we set $\alpha = 1$ for the evaluation of our results. We discuss the implications of choosing the alternative $\alpha(t)$ in more detail in the Results and Discussion section.

The structure of eq 6 allows for the implementation of different models for droplet shape relaxation, as long as it progresses exponentially in the limit of small deviations from equilibrium, as in eq 5. For example, from a microscopic perspective, the motion of the contact line is often described by the so-called molecular kinetic theory (MKT).²⁸ This theory describes the motion of the contact line in terms of small jumps over the intrinsically microscopically inhomogeneous surface, driven by thermal fluctuations. It has been shown to predict contact line dynamics in agreement with experiments and molecular simulation.^{28,45–48} MKT relates the velocity da/dt of the contact line to the driving force via the expression

$$\frac{da}{dt} = \frac{2\xi k_B T \exp\left(-\frac{G^*}{k_B T}\right)}{\eta \nu_L} \sinh\left[\frac{\gamma_{\text{LG}} \xi^2}{2k_B T} (\cos \theta_{\text{eq}} - \cos \theta)\right] \quad (10)$$

where ξ denotes the distance between adsorption sites on the surface; $k_B T$ is the usual thermal energy, with k_B the Boltzmann constant and T the absolute temperature; ν_L is the molecular volume of the liquid; and G^* is the surface contribution to the activation free energy of wetting.^{28,46,49,50}

If we translate eq 10 in terms of the time evolution of the cosine of the contact angle, i.e., make use of eq 1, and expand this to linear order for small deformations $\cos \theta - \cos \theta_{\text{eq}}$, we find that the characteristic time scale τ_{rlx} according to molecular kinetic theory must be given by

$$\tau_{\text{rlx}} = \frac{\eta V^{1/3}}{\gamma_{\text{LG}}} \frac{\nu_L \left(\frac{3}{\pi} \frac{(1 + \cos \theta_{\text{eq}}) \sin \theta_{\text{eq}}}{2 - \cos \theta_{\text{eq}} - \cos^2 \theta_{\text{eq}}}\right)^{1/3}}{\xi^3 \exp\left(-\frac{G^*}{k_B T}\right) (2 + \cos \theta_{\text{eq}}) \sin^2 \theta_{\text{eq}}} \quad (11)$$

We see that the functional form of τ_{rlx} of eq 11 is analogous to that of eq 9. This suggests that the characteristic shape relaxation time as predicted by MKT, which is a microscopic theory in origin, to linear order also is a function of macroscopic parameters such as droplet size, viscosity, and surface tension. Parenthetically, we find that a hydrodynamic theory for contact angle dynamics, as described by Voinov and de Ruijter et al., yields an analogous result.^{45,51} If we expand the theory for small deformations $\cos \theta - \cos \theta_{\text{eq}}$, we again find a characteristic time scale τ_{rlx} which is proportional to the fraction $\eta V^{1/3}/\gamma_{\text{LG}}$. This indicates that the scaling of the characteristic relaxation time scale with viscosity, droplet size, and interfacial tension, as described in eq 9, is universal.

This concludes our analysis of the relaxation dynamics of small drops. We next describe how quasi-steady evaporation affects the dynamics of a deposited droplet, presuming that an instantaneous free energy can be defined, in effect presuming a separation of time scales.

Evaporation of the Droplet. We take quasi-stationary, isothermal vapor diffusion into the surrounding gas phase to be the governing mechanism for evaporation, assuming the droplet to be in contact with an infinite volume of gas. We neglect thermal effects caused by the evaporation of the fluid, effectively assuming that heat transport occurs at much shorter times than the time scales associated with the evaporation process. For water in air, the evaporative cooling at the droplet surface has a negligible effect on the evaporation rate⁵² and we consider an isothermal substrate, which is reasonable for surfaces with high thermal conductivity.^{53–55} Picknett and Bexon¹⁶ derived an expression for the rate of change in mass of a droplet as a function of contact angle θ . The rate of change of the volume V of a droplet can then be written as

$$\frac{dV}{dt} = -\frac{2\pi a D \Delta c}{\rho \sin \theta} f(\theta) \quad (12)$$

where D denotes the diffusion coefficient of vapor molecules in the gas phase and ρ is the mass density of the liquid.¹⁷ Furthermore, $\Delta c \equiv c_s - c_\infty$ denotes the difference between the vapor mass concentration c_s near the liquid–gas interface (in units of mass per volume), presumed to be the saturation value of the fluid molecules in the gas phase, and the vapor mass concentration c_∞ at infinity, i.e., that of the ambient atmosphere. Finally, $f(\theta)$ denotes a geometric factor for which an exact analytical expression is not available in closed form.^{16,56} For our purposes, a polynomial representation for $f(\theta)$

$$f(\theta) = \begin{cases} 0.6366\theta + 0.09591\theta^2 - 0.06144\theta^3 & \text{for } 0 \leq \theta < 0.175 \\ 0.00008957 + 0.6333\theta + 0.116\theta^2 - 0.08878\theta^3 + 0.01033\theta^4 & \text{for } 0.175 \leq \theta \leq \pi \end{cases} \quad (13)$$

is sufficiently accurate. Indeed, the error of the approximant is less than 0.2% for all values of θ .¹⁶

For a constant contact angle θ , eq 12 can be expressed entirely in the contact area radius $a(t)$ using eq 1 and solved exactly. This gives

$$a(t) = \sqrt{a_0^2 - \frac{4D\Delta c}{\rho} \frac{f(\theta) \sin^2 \theta}{2 - 3 \cos \theta + \cos^3 \theta} t} \quad (14)$$

where a_0 denotes the initial value of the contact area radius $a_0 = a(0)$. It shows that the contact area πa^2 decreases linearly in time, a known experimental result.¹⁷ From eq 14, we deduce that the time t_{evap} it takes to evaporate a droplet is the longest for $\theta = \pi/2$. For this contact angle, the evaporation time τ_{evap} is given by the simple expression

$$\tau_{\text{evap}} = \frac{\rho}{2D\Delta c} \left(\frac{3V_0}{2\pi} \right)^{2/3} \quad (15)$$

In the remainder of this work, we shall scale all evaporation times to τ_{evap} . Note that we have identified the two fundamental time scales that describe our problem: the fundamental relaxation time τ_{rlx} (eq 9) and the fundamental evaporation time τ_{evap} (eq 15). The actual evaporation time depends not only on the initial contact angle and the relaxation dynamics of the droplet shape, but also on whether or not contact line pinning takes place.

Contact Line Pinning. Contact line pinning is the phenomenon where the contact line of the droplet becomes stuck, permanently or temporarily, on structural or chemical inhomogeneities of the supporting surface.^{40,57–63} In general, a droplet in the pinned state exhibits a contact angle different from the equilibrium angle θ_{eq} , as it cannot relax to its equilibrium shape. We model the influence of surface heterogeneities by introducing a net macroscopic threshold force per unit length, f_p , exerted in the plane of the surface along the radial direction of the circular contact line. It has a direction opposite to the capillary driving force per unit length, f_c . As both f_p and f_c are exerted on the perimeter of the contact area, we for simplicity refer to both as a force.

If the magnitude of the capillary driving force is smaller than the threshold f_p , then the contact line remains pinned. On the other hand, if it is greater, we allow the contact line to move: the relaxation of the contact angle θ is calculated using eqs 4 and 6 and the contact line moves accordingly. In the presence of contact line pinning, the droplet shape relaxes to the point where the capillary forces and pinning forces are balanced. The contact line motion is quasi-steady and hence the associated friction does not depend on the velocity of the contact line. For simplicity, we presume that the yield force f_p does not depend on the position on the surface. We define the capillary force as

$$f_c = -\frac{1}{2\pi a} \frac{dF}{da} = -\gamma_{\text{LG}}(\cos \theta - \cos \theta_{\text{eq}}) \quad (16)$$

where we have used Young's law (eq 3). In our prescription, we allow motion of the contact line as long as $|f_c| > f_p$. Equation 16 is also referred to as the unbalanced Young's force or unbalanced capillary forces.^{25,33,50}

The magnitude of the pinning force f_p defines a contact angle range in which the capillary force f_c is too weak to overcome pinning. As long as the contact angle θ resides within this range, the contact area remains constant. We refer to this range as the "fixed-area" regime, and it turns out to be bounded by the receding and advancing contact angles, θ_r and θ_a , which are the contact angles for which f_c and f_p are balanced. Within our model, the values of these quantities depend on the pinning force f_p ⁴⁰ according to

$$\theta_r = \arccos(\cos \theta_{\text{eq}} + f_p / \gamma_{\text{LG}}) \quad (17)$$

$$\theta_a = \arccos(\cos \theta_{\text{eq}} - f_p / \gamma_{\text{LG}}) \quad (18)$$

The receding and advancing contact angles indicate the points at which the pinning–depinning transitions occur. If the droplet evaporates while initially being in the pinned (fixed-area) state, the contact angle decreases until the droplet depins at a value equal to θ_r , after which the evaporation continues with a constant contact angle θ , and a receding contact line. In contrast to a constant θ_r , a constant advancing angle θ_a is not encountered for droplets with decreasing volume, but it can only be observed as the point at which the droplet becomes pinned after initial spreading.

RESULTS AND DISCUSSION

We now compare predictions of our phenomenological model with the full nonlinear response presented by molecular kinetic theory (MKT) and with experiments on droplet evaporation in the presence of contact line pinning. We quantify the competition between evaporation and relaxation using the ratio of the two time scales τ_{evap} and τ_{rlx} . It determines, together with the initial and equilibrium contact angles as well as the magnitude of the pinning force, the lifetime of an evaporating droplet. Both fundamental time scales depend only on the properties of the fluid and the surrounding vapor phase. Their ratio scales linearly with the droplet size $V_0^{1/3}$. Presuming that the Arrhenius factor k in eq 9 is of the order 10^8 ,⁴¹ typical values of $\tau_{\text{evap}}/\tau_{\text{rlx}}$ for water droplets of micrometer to millimeter sizes range from 10^{-4} to 10^0 ; however, the latter value may increase further under conditions of slow evaporation (i.e., high humidity). For fluids with higher viscosity η , presuming the other parameters remain unchanged, $\tau_{\text{evap}}/\tau_{\text{rlx}}$ decreases. The ratio of the two time scales has also been addressed by Man and Doi¹⁹ to be important in the context of evaporation problems. Directly connecting the parameter k_{ev} presented in ref 19 to our $\tau_{\text{evap}}/\tau_{\text{rlx}}$ however, is not straightforward due to the factor k . We choose the droplets to be hemispherical in equilibrium, i.e., $\theta_{\text{eq}} = \pi/2$, which is typical for a water droplet on a polymer substrate. The implications of choosing a different equilibrium contact angle are discussed at the end of this section.

Because sessile droplet shape relaxation and evaporation have been described separately in the literature before, we feel it instructive to first investigate how our model compares to those works and to known experimental data. After the validation of the model with the literature, we discuss the predictions given by our more complete model that unites shape relaxation, droplet evaporation, and contact line pinning. Finally, we discuss the impact of the assumptions we make during our calculations.

Shape Relaxation and Pinning–Depinning Transition. To illustrate the relaxation dynamics predicted by our free-energy-based model and to compare the predictions to an existing model for contact line dynamics, we first compare our theory with the relaxation dynamics of a droplet deposited on a surface according to molecular kinetic theory (MKT). This theory, which has a microscopic basis, is shown to describe experimentally measured contact line dynamics rather well.^{28,45–47} As discussed in the Theory section, for small values of $\cos \theta - \cos \theta_{\text{eq}}$, MKT predicts an exponential relaxation with a time scale τ_{rlx} given by eq 11. For greater values, however, the dynamics deviates from a simple single exponential description. To compare the nonlinear contact angle dynamics predicted by our model to that described by MKT, we solve eq 10 numerically. For convenience, we set the equilibrium contact angle to $\theta_{\text{eq}} = \pi/2$ and choose four initial

angles θ_0 symmetrically around this angle. In Figure 2, we compare the time dependence of the contact angle θ and the

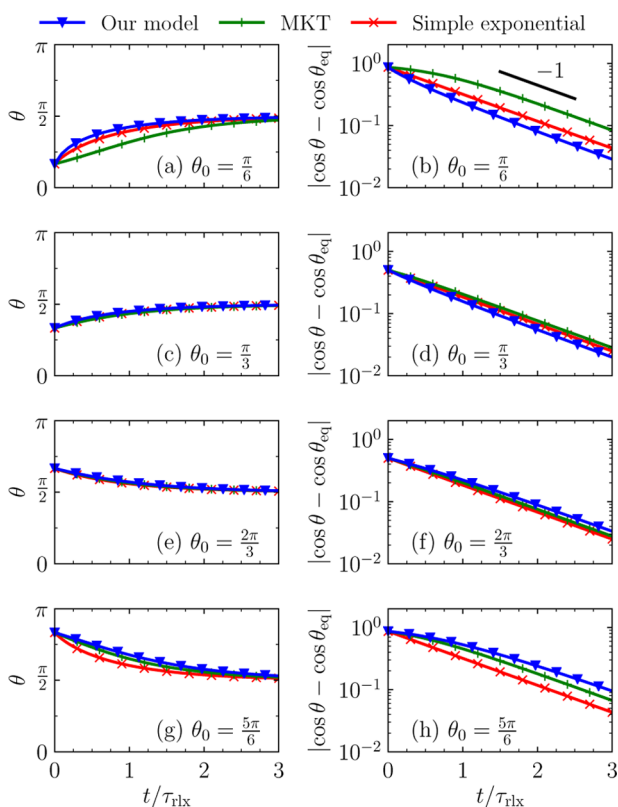


Figure 2. Comparison between predictions for the relaxation dynamics of the contact angle θ to its equilibrium value $\theta_{\text{eq}} = \pi/2$ from our model (blue triangles), molecular kinetic theory (green pluses), and a simple exponential relaxation for comparison (red crosses). Left: contact angle θ as a function of dimensionless time t/τ_{rlx} ; right: the corresponding $|\cos \theta - \cos \theta_{\text{eq}}|$.

absolute value of the difference between the cosines of $\theta(t)$ and θ_{eq} . Indicated in the figures are the results of our model (blue triangles), the MKT result (green pluses), and a simple single exponential relaxation as given in eq 5 (red crosses). Note that, as in all cases τ_{rlx} and θ_{eq} are fixed, the results indicated in the figures as “our model” are independent of the choice of the characteristic time scale and hence also describe the result for, e.g., the linearized version of the MKT model.

Figure 2 informs us of the following:

- For deviations of $\pm\pi/6$ from the equilibrium value of $\pi/2$ (see Figure 2c,e), the agreement between the evolution of the contact angle as a function of scaled time predicted by all three descriptions is excellent. For greater initial deviations from the equilibrium angle (Figure 2a,g), the agreement remains remarkably good, in particular for the larger initial angle.
- Figure 2d,f highlights any inconsistencies for small deviations from the equilibrium by focusing on the difference of the cosines on a logarithmic scale. These figures show that well within one characteristic time scale simple single exponential decay is reached. Any small late-stage deviations between the curves is caused by the early-stage nonlinear behavior. Figure 2b,h shows that even for greater initial deviations from the

equilibrium contact angle, simple single exponential decay occurs within one characteristic time scale.

The process of droplet evaporation in the presence of contact line pinning has been studied theoretically by Stauber et al.,²⁴ who describe the dependence of the evaporation time t_{evap} on the initial contact angle θ_0 , where they fix the receding contact angle θ_r to several values. They consider two separate modes of evaporation, a constant contact radius (CCR, pinned) and a constant contact angle (CCA, receding) mode, allow for pinning–depinning transitions and model the evaporation dynamics accordingly using an evaporation description analogous to eq 12. Their results can be reproduced quantitatively by our model. However, our model also includes the relaxation of the droplet shape toward its equilibrium angle, after it is deposited on the surface with an angle different from the equilibrium value. We discuss in more detail the similarities and differences between their work and the results from our model in the next subsection.

We now relate results from our model to the experimental data of Belmiloud et al.⁶ on the evaporation of a water droplet on a flat silicon surface (see Figure 3). Figure 3 shows the

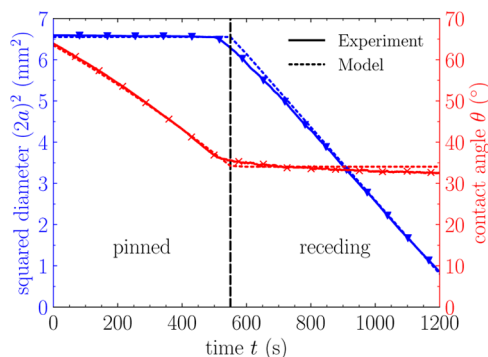


Figure 3. Comparison between the results of experiments on the evaporation of sessile water droplets on a silica wafer⁶ (solid lines) and the numerical evaluation of the droplet model (dashed lines). The squared contact diameter $(2a)^2$ (left vertical axis, blue triangles) and contact angle θ (right vertical axis, red crosses) are shown as a function of time t . The two modes of evaporation, pinned and receding, are indicated. The model parameters are as follows: pinning force, $f_p \approx 0.034 \text{ N m}^{-1}$; vapor concentration difference, $\Delta c = 11.6 \times 10^{-3} \text{ kg m}^{-3}$; and vapor diffusion coefficient, $D = 3.15 \times 10^{-5} \text{ m}^2 \text{ s}^{-1}$.

squared contact diameter $(2a)^2$ (blue triangles) and contact angle θ (red crosses) as a function of time t . The results of Belmiloud et al., represented by the solid lines, can be readily reproduced by our phenomenological model (dashed lines). Initially, the contact line of the droplet is pinned, as is seen from the squared contact diameter remaining constant, while the contact angle decreases. As the receding contact angle θ_r is reached, a pinning–depinning transition occurs, after which the angle remains constant at the receding value and the diameter squared decreases linearly, as discussed in the Theory section.

To model the evaporation process of the initially pinned droplet, we choose our model parameters to correspond to the experimental values. The pinning force per unit length f_p was set to a value of $f_p \approx 0.034 \text{ N m}^{-1}$ to obtain a receding contact angle of $\theta_r = 34^\circ$. The value of the pinning force per unit length is of the same order of magnitude as the liquid–air interfacial tension ($\gamma_{\text{LG}} = 0.07 \text{ N m}^{-1}$). The values reported by Belmiloud et al.⁶ for the surface vapor concentration c_s and the

relative humidity were used to determine $\Delta c = 11.6 \times 10^{-3} \text{ kg m}^{-3}$. The best correspondence between the measurement and our model is obtained for a vapor diffusion coefficient $D = 3.15 \times 10^{-5} \text{ m}^2 \text{ s}^{-1}$, as opposed to the reported $D = 2.60 \times 10^{-5} \text{ m}^2 \text{ s}^{-1}$. However, Belmiloud et al. also report on an underestimation of the evaporation rate: the droplet evaporates faster than predicted by eq 12.⁶ This is arguably due to inaccuracies in measuring the properties of the ambient vapor.

Predictions by Full Model. We now consider the effect of the interplay between the three components of our model to predict the evaporation dynamics of a droplet. To that end, we first discuss two limiting cases. We report our findings on (1) the effect of contact line pinning on a nonevaporating, relaxing droplet and (2) the effect of shape relaxation on the lifetimes of droplets with an unpinned contact line. Subsequently, we present our results on simultaneous shape relaxation and evaporation of a droplet subject to contact line pinning.

If the shape relaxation of a droplet is affected by contact line pinning, the contact angle relaxation in the absence of evaporation studied in the previous section (Figure 2) changes drastically, as is illustrated in Figure 4. If droplets start out

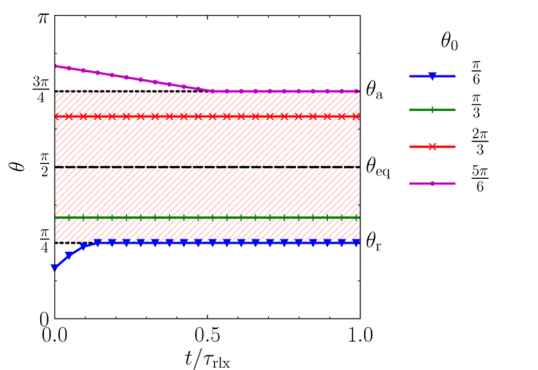


Figure 4. Relaxation of the contact angle θ of a deposited drop on a surface toward the equilibrium value $\theta_{\text{eq}} = \pi/2$. There is no evaporation and the pinning force f_p is set such that the receding and advancing contact angles are $\theta_r = \pi/4$ and $\theta_a = 3\pi/4$. This results in the fixed-area range between θ_r and θ_a , where the capillary force f_c cannot overcome f_p and the contact line becomes or remains pinned.

within the fixed-area region, i.e., have an initial angle $\theta_r < \theta_0 < \theta_a$, indicated by the shaded region in Figure 4, then the contact line is not able to move. In other words, the droplets are not able to relax their shape to accommodate the equilibrium contact angle θ_{eq} . For initial angles outside of this regime, shape relaxation does occur, albeit only until the fixed-area region is reached, after which the motion of the contact line is halted. This phenomenon has strong implications for the lifetime of an evaporating droplet. The asymmetry in the time it takes for the droplet to become pinned for $\theta_0 = \pi/6$ and for $\theta_0 = 5\pi/6$ has its origin again, as is the case for the shape relaxation shown in Figure 2, in the nonlinearity of eqs 4 and 6. We note that the curves shown in Figure 4 depict the relaxation of the contact angle θ . The exponential relaxation of the cosine in eq 5 is therefore not immediately evident from the figure.

If we allow for evaporation, the shape relaxation of a droplet from an initial contact angle θ_0 toward its equilibrium angle θ_{eq} may have a strong impact on the evaporation dynamics of a droplet, also without any contact line pinning occurring. As discussed in the Theory section, the evaporation rate depends

on the contact angle θ , and is at its minimum for $\theta = \pi/2$. If a droplet with a certain θ_{eq} is deposited onto a surface with an initial angle $\theta_0 \neq \theta_{\text{eq}}$, the relative speeds at which the droplet relaxes to its equilibrium angle and at which it evaporates, characterized by the ratio $\tau_{\text{evap}}/\tau_{\text{rlx}}$, will influence the lifetime of such a droplet. In the remainder of our manuscript, we adopt the representation style of Stauber et al.²⁴ when discussing lifetimes of droplets, where we depict the scaled evaporation time $t_{\text{evap}}/\tau_{\text{evap}}$ as a function of the initial contact angle θ_0 .

First, we consider the evaporation of droplets in the absence of contact line pinning. In Figure 5, we present the droplet

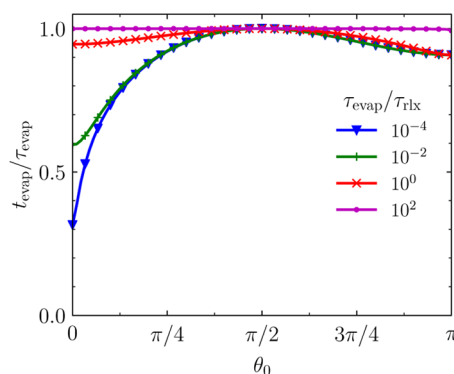


Figure 5. Scaled evaporation times $t_{\text{evap}}/\tau_{\text{evap}}$ for sessile droplets as a function of the initial contact angle θ_0 , for ratios $\tau_{\text{evap}}/\tau_{\text{rlx}} = 10^{-4}, 10^{-2}, 10^0$, and 10^2 . The equilibrium contact angle is $\theta_{\text{eq}} = \pi/2$.

lifetimes $t_{\text{evap}}/\tau_{\text{evap}}$ as a function of the initial contact angle θ_0 , for $\tau_{\text{evap}}/\tau_{\text{rlx}} = 10^{-4}, 10^{-2}, 10^0$, and 10^2 . In the limit of slow shape relaxation ($\tau_{\text{evap}}/\tau_{\text{rlx}} \ll 1$), we exactly recover the result of Stauber et al.²⁴ for evaporation with a constant contact angle (see the blue triangles in Figure 5). The droplet lifetime decreases rapidly for $\theta_0 \rightarrow 0$, as the area-to-volume ratio increases. As already discussed in the Theory section, the lifetime is longest for $\theta = \pi/2$, resulting in a maximum in the graph. For $\theta > \pi/2$, the lifetimes slightly decrease again due to the increase in the area-to-volume ratio.

For increasing $\tau_{\text{evap}}/\tau_{\text{rlx}}$ we see a decreasing effect of the initial angle on the lifetimes, as the contact angles relax more quickly to the equilibrium value $\theta_{\text{eq}} = \pi/2$. The increase in droplet lifetime due to faster shape relaxation is most notable for small contact angles, as the relaxation is the fastest in that regime and small changes to the contact angle induce large changes in the evaporation rate, according to eq 12. Close to $\theta_0 = \pi/2$, however, both shape relaxation and evaporation are slow and hence the changes in droplet lifetime upon changing $\tau_{\text{evap}}/\tau_{\text{rlx}}$ are small. For $\tau_{\text{evap}}/\tau_{\text{rlx}} \gg 1$, the droplet lifetime is effectively independent of θ_0 . We conclude that the evaporation dynamics of a droplet on a flat surface is strongly affected by the ratio between the rates of evaporation and shape relaxation, in the absence of contact line pinning.

If the contact line can become stuck on the surface due to pinning, however, the response of a drop to deposition on a surface becomes more complex. We can identify three regimes in the droplet dynamics, distinguished by the value of the initial contact angle θ_0 relative to the receding and advancing contact angles θ_r and θ_a , respectively

- $\theta_0 < \theta_r$, where θ_0 lies below the fixed-area range;
- $\theta_r \leq \theta_0 \leq \theta_a$, where θ_0 lies within the fixed-area range;
- $\theta_0 > \theta_a$, where θ_0 lies above the fixed-area range.

Figure 6 shows the droplet shape in terms of the scaled squared radius $(a/a_0)^2$ (blue triangles) and θ (red crosses) as a

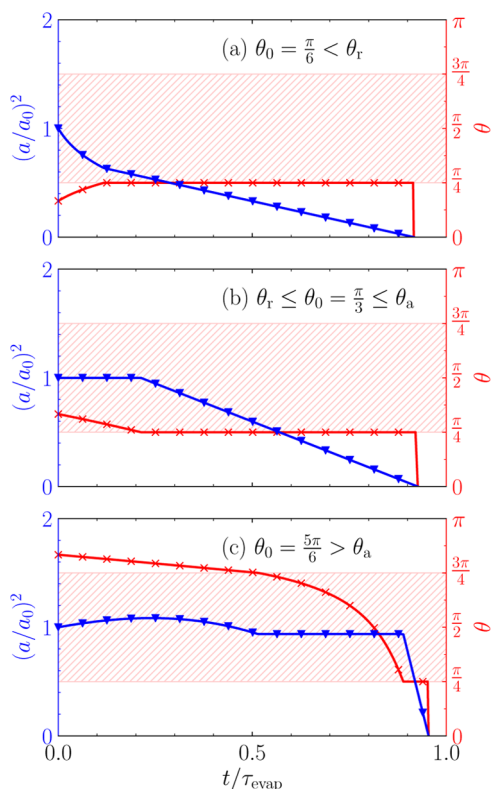


Figure 6. Scaled contact area $(a/a_0)^2$ and contact angle θ as a function of nondimensional time t/τ_{evap} , for the evaporation of a droplet with equilibrium angle $\theta_{\text{eq}} = \pi/2$. The shaded areas indicate the fixed-area domains between the receding and advancing contact angles, $\theta_r = \pi/4$ and $\theta_a = 3\pi/4$, respectively. The ratio between evaporation and relaxation time scales is set to $\tau_{\text{evap}}/\tau_{\text{rlx}} = 1$, and we show the time evolution of the droplet shape for three values of the initial contact angle θ_0 : (a) $\theta_0 = \pi/6 < \theta_r$, (b) $\theta_r < \theta_0 = \pi/3 < \theta_a$, and (c) $\theta_0 = 5\pi/6 > \theta_a$.

function of nondimensional time t/τ_{evap} . The pinning force f_p is set to $f_p \approx 0.924 \gamma_{\text{LG}}$, leading to $\theta_r = \pi/4$ and $\theta_a = 3\pi/4$. The contact angle range between the two (the fixed-area domain) is indicated by the shaded area. The ratio between evaporation and relaxation time scales is $\tau_{\text{evap}}/\tau_{\text{rlx}} = 1$. Three different graphs are shown to illustrate the three regimes. If $\theta_0 < \theta_r$, the droplet starts out in the depinned state and thus the contact line moves to accommodate the contact angle relaxation toward the equilibrium value θ_{eq} (see Figure 6a). After some time, the receding contact angle θ_r is reached and the contact angle relaxation halts, resulting in the remainder of the evaporation process occurring with a constant contact angle θ_r . In the second case, shown in Figure 6b, the droplet starts out with a pinned contact line, since $\theta_r < \theta_0 < \theta_a$. Due to evaporation, the contact angle decreases until it reaches θ_r . At that point, a depinning transition occurs, the contact line is allowed to move again and evaporation continues with a constant contact angle θ_r . Parenthetically, in Figure 6b, we present the results for an initial contact angle $\theta_0 < \pi/2$, but the general behavior of the contact angle as a function of time is the same for $\pi/2 \leq \theta_0 < \theta_a$. Finally, for $\theta_0 > \theta_a$, the contact line can initially move freely, causing the droplet to spread and increase its contact area (see Figure 6c). After the contact

angle reaches the advancing value θ_a , however, the contact line becomes pinned. Once more, from this point on, the contact angle decreases due to evaporation. And again, as the contact angle reaches the receding value θ_r , a depinning transition occurs and evaporation continues with a constant contact angle.

The emergence of the three regimes due to the presence of contact line pinning has a significant impact on the droplet lifetimes. The extent of the effect depends on the ratio of evaporation to relaxation time scales $\tau_{\text{evap}}/\tau_{\text{rlx}}$, as illustrated in Figure 7. Figure 7 shows the scaled lifetime of an evaporating

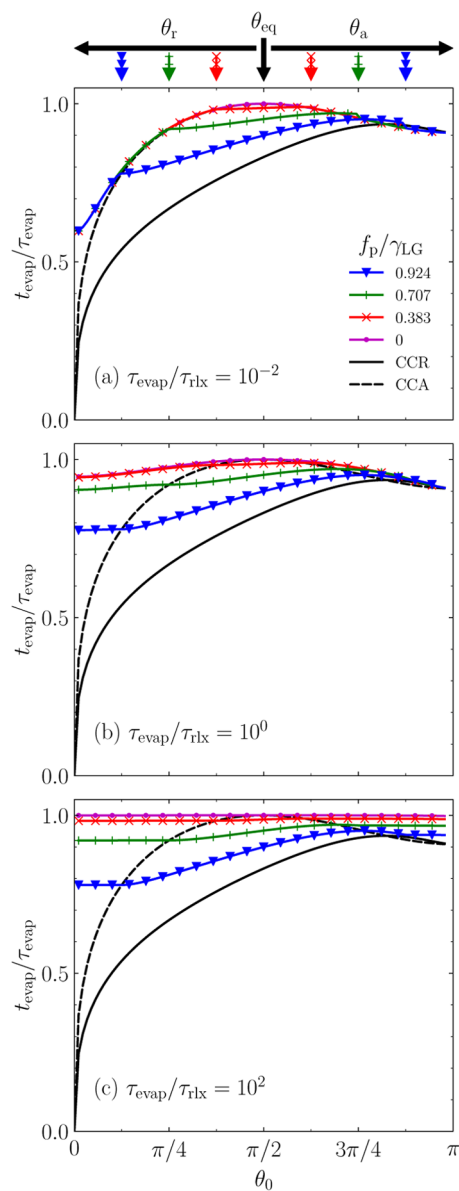


Figure 7. Scaled evaporation times $t_{\text{evap}}/\tau_{\text{evap}}$ of sessile droplets as a function of the initial contact angle θ_0 , for various values for $f_p/\gamma_{\text{LG}} \approx 0.924, 0.707, 0.383$, and 0 , and three ratios $\tau_{\text{evap}}/\tau_{\text{rlx}}$ of (a) 10^{-2} , (b) 10^0 , and (c) 10^2 , covering the range from fast to slow evaporation. The values of the respective receding and advancing contact angles are indicated by arrows in the same color as the corresponding lines in the graph. The black lines represent the two limiting cases of evaporation at constant radius of the contact area (CCR, solid) and evaporation at a constant contact angle (CCA, dashed). The equilibrium contact angle is $\theta_{\text{eq}} = \pi/2$.

droplet $t_{\text{evap}}/\tau_{\text{evap}}$ as a function of the initial contact angle θ_0 , for $\tau_{\text{evap}}/\tau_{\text{rlx}} = 10^{-2}$, 10^0 , and 10^2 , covering the entire range from fast to slow evaporation. We can identify two limiting cases for the evaporation in all three graphs, shown in black, being evaporation with a constant contact radius (CCR, solid lines) and evaporation with a constant contact angle (CCA, dashed lines). These two limits are not dependent on the ratio $\tau_{\text{evap}}/\tau_{\text{rlx}}$ as we impose that either the contact area or the constant angle remains fixed.

The time it takes for a droplet to evaporate is shorter at a constant radius compared to a constant angle for the majority of the initial angle range $0 \leq \theta_0 \leq \pi$. This is because the constant contact radius mode causes the contact angle to decrease during evaporation. Decreasing θ generally speeds up the evaporation process due to an increase of the surface-to-volume ratio, especially at late times. For large initial angles ($\theta_0 \rightarrow \pi$), however, evaporation in the constant-angle mode becomes faster than evaporation in the constant-radius mode. As the latter causes a continuous decrease in the contact angle, it initially slows down the evaporation rate before speeding it up again. As discussed before, the maximum lifetime of a droplet evaporating in the constant-angle mode is $t_{\text{evap}} = \tau_{\text{evap}}$ for $\theta_0 = \pi/2$, resulting in a maximum in the graph. For the constant-radius mode, the maximum lifetime is shorter than the maximum in the constant-angle mode, being $t_{\text{evap}} \approx 0.9354\tau_{\text{evap}}$ for $\theta_0 \approx 0.822\pi$.²⁴

The differently colored arrows at the top of Figure 7 represent the receding contact angle θ_r and the advancing contact angle θ_a . These define the domain in which the contact area remains constant. The arrow colors correspond to the colors of the curves shown in the figure, which depict the droplet lifetimes $t_{\text{evap}}/\tau_{\text{evap}}$ as a function of the initial contact angle θ_0 , for values of the pinning force f_p/γ_{LG} of approximately 0.924 (blue triangles), 0.707 (green pluses), 0.383 (red crosses), and 0 (purple dots). The values for θ_r and θ_a remain constant in all three graphs, and as $\theta_{\text{eq}} = \pi/2$, they take symmetric values around the equilibrium.

For all three graphs in Figure 7a–c, the curve segments between the bounding receding and advancing contact angles are identical. The reason is that if the contact line of a drop is initially pinned due to the choice of initial angle, this angle cannot relax toward its equilibrium value as is also shown in Figure 6b. Therefore, the magnitude of the shape relaxation rate does not affect the evaporation process. During evaporation, the contact angle only decreases from the initial to the receding angle and remains at that value until the drop has fully evaporated.

The relaxation of the contact angle toward its equilibrium value is only possible for initial angles outside of the fixed-area domain, as shown in Figure 6a,c, where shape relaxation occurs until the contact angle reaches either boundary. In other words, for values of θ_0 outside of the fixed-area region, the relative shape relaxation rate does have an impact on the droplet lifetime. For values of $\tau_{\text{evap}}/\tau_{\text{rlx}} \gtrsim 1$, as shown in Figure 7b,c, evaporation is relatively slow and relaxation, in essence, is instantaneous. This leads to an evaporation time that is essentially an invariant of the initial angle, outside of the fixed-area domain, where the lifetime takes on the value at the nearest boundary (at θ_r or θ_a). If evaporation is very fast, i.e., $\tau_{\text{evap}}/\tau_{\text{rlx}} \ll 1$, as shown in Figure 7a, relaxation cannot keep up and the evaporation time is dictated by a virtually constant contact angle. For sufficiently small θ_0 , however, relaxation can keep up with evaporation and the evaporation time deviates

from the lifetimes for the constant contact angle mode. This deviation vanishes for $\tau_{\text{evap}}/\tau_{\text{rlx}} \rightarrow 0$.

For initial angles above the advancing angle, the lifetime curves start to deviate from both limiting cases and from the curves reported by Stauber et al.,²⁴ when $\tau_{\text{evap}}/\tau_{\text{rlx}}$ increases. This is caused by the evaporation dynamics predicted by our model being more complicated than a simple imposed transition from a pinned into a depinned state. As shown in Figure 6c, the contact line can initially move freely, implying that the contact angle starts to move toward its equilibrium value θ_{eq} . Upon reaching θ_a , the contact line becomes pinned and the contact angle decreases until it reaches θ_r . Subsequently, a pinning–depinning transition occurs and the droplet evaporates with a fixed contact angle. In other words, the droplet experiences two transitions, rather than one, by subsequently going through depinned, pinned, and depinned modes.

Impact of Assumptions. We developed a phenomenological model for the shape relaxation of an evaporating droplet. The characteristic time scale associated with this relaxation is found to be proportional to a length scale L . This length scale L has been connected to (1) a slip or friction length or (2) the size of the droplet. To incorporate the effects of either length scale on the evaporation dynamics, we have equipped eq 6, which describes the droplet relaxation, with a scale factor α . In the discussion of the results above, we chose $\alpha = 1$ for simplicity.

Now we discuss in more detail the implications of considering an alternative $\alpha = \alpha(t)$, which is proportional to the droplet size

$$\alpha(t) = \left[\frac{V(t)}{V_0} \right]^{1/3} \quad (19)$$

where $V(t)$ and V_0 denote the instantaneous and initial droplet volumes, respectively. As a consequence, the relaxation process speeds up as the droplet size decreases. We find, however, that explicitly taking this effect into account hardly affects the droplet lifetime. This is caused by the circumstance that the capillary driving force is the strongest at short times, as the difference $\cos\theta - \cos\theta_{\text{eq}}$ is then the greatest. In other words, the majority of the relaxation process occurs at short times. However, at short times, the droplet has hardly lost any volume by evaporation, which means that the scale factor $\alpha(t) \approx 1$, causing the relaxation processes for both expressions for α to occur in virtually the same manner. In Figure 8, we depict the squared scaled radius $(a/a_0)^2$ (blue triangles) and the contact angle θ (red crosses) of evaporating droplets as a function of scaled time t/τ_{evap} . We present the results for the scale factor $\alpha(t) = [V(t)/V_0]^{1/3}$ (solid lines) compared to $\alpha = 1$ (dashed lines), for three values of θ_0 . For these calculations, we do not incorporate a contact line pinning force, which means the droplet is allowed to relax its shape toward $\theta_{\text{eq}} = \pi/2$, and $\tau_{\text{evap}}/\tau_{\text{rlx}} = 1$.

In all three graphs, we can clearly see that the dynamics described by the two expressions for α are identical at early times. Only after the droplet has partly evaporated, we see a slight deviation in the dynamics, due to the decrease of $\alpha(t)$. This effect only arises after approximately 30% of the evaporation time has passed. For large initial contact angles θ_0 , which we show in Figure 8c, the deviations between the graphs for the two expressions for α are slightly larger than those for smaller θ_0 (Figure 8a,b). However, the time at which

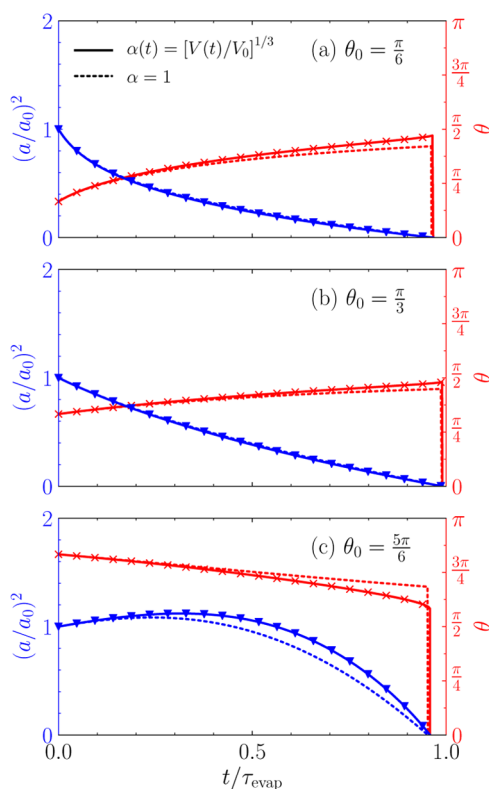


Figure 8. Comparison of the squared scaled radius $(a/a_0)^2$ and the contact angle θ between the alternative scale factor $\alpha(t) = [V(t)/V_0]^{1/3}$ (solid lines) and the original $\alpha = 1$ (dashed lines), as a function of scaled time t/τ_{evap} . $\tau_{\text{evap}}/\tau_{\text{rlx}} = 1$ and figures are shown for initial angles: (a) $\theta_0 = \pi/6$, (b) $\theta_0 = \pi/3$, (c) $\theta_0 = 5\pi/6$.

the droplet is fully evaporated is hardly affected. Note that the presence of contact line pinning would only decrease the effect shape relaxation has on the evaporation time, as it inhibits contact line motion for a certain range of the contact angle θ . We conclude from this that the lifetime of an evaporating droplet is not sensitive to our choice of α , hence explicitly taking into account the size dependence of the relaxation process has a negligible effect on the total evaporation time. We note that this phenomenon is valid regardless of the value of $\tau_{\text{evap}}/\tau_{\text{rlx}}$: a reproduction of Figure 7 with the alternative scale factor α yields the same graph.

A second point left for discussion is the influence of the value of the equilibrium angle θ_{eq} on the evaporation dynamics of a droplet. We recall that an equilibrium contact angle of $\pi/2$ has two implications for the evaporation dynamics of a droplet. First, evaporation is slowest at $\theta = \pi/2$, as described by eq 12. Second, in the presence of contact line pinning, the fixed-area domain is located symmetrically around the equilibrium angle: as $\cos \theta_{\text{eq}} = 0$, the advancing contact angle is given by $\theta_a = \pi - \theta_r$, as described by eqs 17 and 18. Both properties are not valid for droplets with an equilibrium contact angle $\theta_{\text{eq}} \neq \pi/2$. A droplet taking on a nonhemispherical equilibrium results, on the one hand, in the fact that faster shape relaxation, or increasing $\tau_{\text{evap}}/\tau_{\text{rlx}}$, not necessarily implies slower evaporation, as we have shown in Figure 7, but that this depends on the initial and equilibrium contact angles. On the other hand, the receding and advancing contact angles are now located asymmetrically around the equilibrium angle.

As we have seen above, if contact line pinning occurs, the time it takes for a droplet to evaporate depends strongly on the

values of the receding and advancing contact angles (see Figure 7). For contact angles in between the two, contact line motion is inhibited. For $\theta_{\text{eq}} \neq \pi/2$, this principle is still valid, only θ_r and θ_a are located asymmetrically around θ_{eq} . For initial angles within the fixed-area domain, the lifetimes remain unchanged with respect to $\theta_{\text{eq}} = \pi/2$. As contact line motion is inhibited there, shape relaxation is blocked, so the value of θ_{eq} is irrelevant. For initial angles outside of the domain, the lifetime as a function of initial angle behaves similar to what we show in Figure 7: for increasing shape relaxation rates, i.e., increasing $\tau_{\text{evap}}/\tau_{\text{rlx}}$ the evaporation time of a droplet converges to the values at the boundaries and it becomes increasingly less dependent on the initial contact angle θ_0 .

We conclude from the above that the specific value of the equilibrium contact angle θ_{eq} has little effect on the general behavior of the droplet lifetime as a function of the initial contact angle. It does affect the evaporation dynamics, but to an extent that is limited to two factors. On the one hand, it determines the evaporation time at the equilibrium angle, so it affects the droplet lifetime most in the absence of contact line pinning and in the limit of fast shape relaxation. On the other hand, it determines, together with the magnitude of the pinning force f_p , the locations of the receding and advancing contact angles, which in turn define the region in which contact line motion is inhibited.

■ SUMMARY AND CONCLUSIONS

In conclusion, we propose a model for diffusive evaporation of a droplet on a flat surface, which accounts for the relaxation of the contact angle toward its equilibrium value. This shape relaxation is driven by the tendency of the droplet to reach its minimum free-energy state. We also model pinning of the contact line onto the surface by introducing a pinning force, insisting that the contact line remains pinned as long as the capillary forces are not able to overcome this threshold force.

Within our model description, the time it takes for a droplet to evaporate turns out to depend on five parameters: the initial and equilibrium contact angles, the characteristic time scales associated with shape relaxation and evaporation, and the magnitude of the contact line pinning force. The ratio between the two characteristic time scales describes the competition between shape relaxation and evaporation, which has a significant effect on the droplets' lifetime. In the limit of slow relaxation (or fast evaporation), the total evaporation time of a droplet strongly depends on the initial contact angle, whereas for fast relaxation, the lifetime is virtually unaffected by the value of the initial contact angle.

The presence of a pinning force results in a contact angle range for which the contact line is fixed, as the capillary forces are not capable of overcoming the pinning force. This regime is bounded by the receding and advancing contact angles and as long as the contact angle resides within this range, the contact area remains constant. The magnitude of the pinning force determines the values of the receding and advancing contact angles and therefore has an impact on the lifetime of an evaporating droplet: the shape relaxation of a droplet becomes partly suppressed because the droplet cannot relax its shape for contact angles within this fixed-area regime.

We show that shape relaxation has a significant impact on the evaporation time of a droplet, both in the absence and in the presence of contact line pinning. Explicitly taking into account the size dependence of the relaxation process turns out to have virtually no effect on the droplet's lifetime, since the

majority of the relaxation occurs at short times for which the droplet size has hardly decreased. The value of θ_{eq} does also not affect the general dynamical behavior; however, it does define the lifetime for a droplet at its equilibrium angle and the location of the receding and advancing contact angles.

Finally, the simplicity of our model allows for relatively straightforward evaluation of the dynamics of an evaporating droplet. This means that it can also be readily extended to, e.g., take into account compound exchange between the solid phase and the liquid phase, or investigate an evaporation process wherein the droplet properties do not remain constant in time.

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Notes

The authors declare no competing financial interest.

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REFERENCES

- (1) Kuang, M.; Wang, L.; Song, Y. Controllable Printing Droplets for High-Resolution Patterns. *Adv. Mater.* **2014**, *26*, 6950–6958.
- (2) Park, J.; Moon, J. Control of Colloidal Particle Deposit Patterns within Picoliter Droplets Ejected by Ink-Jet Printing. *Langmuir* **2006**, *22*, 3506–3513.
- (3) Calvert, P. Inkjet Printing for Materials and Devices. *Chem. Mater.* **2001**, *13*, 3299–3305.
- (4) Yu, Y.; Zhu, H.; Frantz, J. M.; Reding, M. E.; Chan, K. C.; Ozkan, H. E. Evaporation and coverage area of pesticide droplets on hairy and waxy leaves. *Biosyst. Eng.* **2009**, *104*, 324–334.
- (5) Wei, Y.; Brainard, R. L. *Advanced Processes for 193-nm Immersion Lithography*; SPIE: Bellingham, 2009.
- (6) Belmiloud, N.; Tamaddon, A. H.; Mertens, P. W.; Struyf, H.; Xu, X. Dynamics of the Drying Defects Left by Residual Ultra-Pure Water Droplets on Silicon Substrate. *ECS J. Solid State Sci. Technol.* **2012**, *1*, P34–P39.
- (7) Mack, C. *Fundamental Principles of Optical Lithography*; John Wiley & Sons, Ltd: Chichester, U.K., 2007.
- (8) Lin, B. J. The future of subhalf-micrometer optical lithography. *Microelectron. Eng.* **1987**, *6*, 31–51.
- (9) Kocsis, M.; van den Heuvel, D.; Gronheid, R.; Maenhoudt, M.; Vangoidsenhoven, D.; Wells, G.; Stepanenko, N.; Benndorf, M.; Kim, H. W.; Kishimura, S.; Ercken, M.; van Roey, F.; O'Brien, S.; Fyen, W.; Foubert, P.; Moerman, R.; Streefkerk, B. In *Immersion Specific Defect Mechanisms: Findings and Recommendations for Their Control*, Proceedings of SPIE, Optical Microlithography XIX, 2006; Vol. 6154, 615409.
- (10) Bourgès-Monnier, C.; Shanahan, M. E. R. Influence of Evaporation on Contact Angle. *Langmuir* **1995**, *11*, 2820–2829.
- (11) Uno, K.; Hayashi, K.; Hayashi, T.; Ito, K.; Kitano, H. Particle adsorption in evaporating droplets of polymer latex dispersions on hydrophilic and hydrophobic surfaces. *Colloid Polym. Sci.* **1998**, *276*, 810–815.
- (12) Cachile, M.; Bénichou, O.; Poulard, C.; Cazabat, A. M. Evaporating Droplets. *Langmuir* **2002**, *18*, 8070–8078.

- (13) Fukai, J.; Ishizuka, H.; Sakai, Y.; Kaneda, M.; Morita, M.; Takahara, A. Effects of droplet size and solute concentration on drying process of polymer solution droplets deposited on homogeneous surfaces. *Int. J. Heat Mass Transfer* **2006**, *49*, 3561–3567.

- (14) Yu, Y.-S.; Wang, Z.; Zhao, Y.-P. Experimental and theoretical investigations of evaporation of sessile water droplet on hydrophobic surfaces. *J. Colloid Interface Sci.* **2012**, *365*, 254–259.

- (15) Hu, H.; Larson, R. G. Evaporation of a Sessile Droplet on a Substrate. *J. Phys. Chem. B* **2002**, *106*, 1334–1344.

- (16) Picknett, R.; Bexon, R. The evaporation of sessile or pendant drops in still air. *J. Colloid Interface Sci.* **1977**, *61*, 336–350.

- (17) Erbil, H. Y.; McHale, G.; Newton, M. I. Drop Evaporation on Solid Surfaces: Constant Contact Angle Mode. *Langmuir* **2002**, *18*, 2636–2641.

- (18) Hughes, A. P.; Thiele, U.; Archer, A. J. Liquid drops on a surface: Using density functional theory to calculate the binding potential and drop profiles and comparing with results from mesoscopic modelling. *J. Chem. Phys.* **2015**, *142*, No. 074702.

- (19) Man, X.; Doi, M. Ring to Mountain Transition in Deposition Pattern of Drying Droplets. *Phys. Rev. Lett.* **2016**, *116*, No. 066101.

- (20) Frank, X.; Perré, P. Droplet spreading on a porous surface: A lattice Boltzmann study. *Phys. Fluids* **2012**, *24*, No. 042101.

- (21) Ledesma-Aguilar, R.; Vella, D.; Yeomans, J. M. Lattice-Boltzmann simulations of droplet evaporation. *Soft Matter* **2014**, *10*, 8267–8275.

- (22) Hirvi, J. T.; Pakkanen, T. A. Molecular dynamics simulations of water droplets on polymer surfaces. *J. Chem. Phys.* **2006**, *125*, No. 144712.

- (23) Zhang, J.; Leroy, F.; Müller-Plathe, F. Influence of Contact-Line Curvature on the Evaporation of Nanodroplets from Solid Substrates. *Phys. Rev. Lett.* **2014**, *113*, No. 046101.

- (24) Stauber, J. M.; Wilson, S. K.; Duffy, B. R.; Sefiane, K. On the lifetimes of evaporating droplets. *J. Fluid Mech.* **2014**, *744*, R2.

- (25) Stauber, J. M.; Wilson, S. K.; Duffy, B. R.; Sefiane, K. On the lifetimes of evaporating droplets with related initial and receding contact angles. *Phys. Fluids* **2015**, *27*, No. 122101.

- (26) Schonhorn, H.; Frisch, H.; Kwei, T. Kinetics of Wetting of Surfaces by Polymer Melts. *J. Appl. Phys.* **1966**, *37*, 4967.

- (27) Newman, S. Kinetics of wetting of surfaces by polymers; capillary flow. *J. Colloid Interface Sci.* **1968**, *26*, 209–213.

- (28) Blake, T.; Haynes, J. Kinetics of displacement. *J. Colloid Interface Sci.* **1969**, *30*, 421–423.

- (29) de Gennes, P.-G.; Brochard-Wyart, F.; Quéré, D. *Capillarity and Wetting Phenomena*; Springer: New York, 2004.

- (30) Extrand, C. W.; Kumagai, Y. Contact Angles and Hysteresis on Soft Surfaces. *J. Colloid Interface Sci.* **1996**, *184*, 191–200.

- (31) Whyman, G.; Bormashenko, E.; Stein, T. The rigorous derivation of Young, Cassie-Baxter and Wenzel equations and the analysis of the contact angle hysteresis phenomenon. *Chem. Phys. Lett.* **2008**, *450*, 355–359.

- (32) Young, T. An Essay on the Cohesion of Fluids. *Philos. Trans. R. Soc. London* **1805**, *95*, 65–87.

- (33) de Gennes, P. G. Wetting: statics and dynamics. *Rev. Mod. Phys.* **1985**, *57*, 827–863.

- (34) Montes Ruiz-Cabello, F. J.; Rodríguez-Valverde, M. A.; Cabrerizo-Vílchez, M. A. Equilibrium contact angle or the most-stable contact angle? *Adv. Colloid Interface Sci.* **2014**, *206*, 320–327.

- (35) Swift, J.; Hohenberg, P. C. Hydrodynamic fluctuations at the convective instability. *Phys. Rev. A* **1977**, *15*, 319–328.

- (36) Goldenfeld, N. D. *Lectures on Phase Transitions and the Renormalization Group*; Perseus Books: New York, 1992; p 209.

- (37) Chaikin, P. M.; Lubensky, T. C. *Principles of Condensed Matter Physics*; Cambridge University Press, 1995; pp 466–468.

- (38) Hårth, M.; Schubert, D. W. Simple Approach for Spreading Dynamics of Polymeric Fluids. *Macromol. Chem. Phys.* **2012**, *213*, 654–665.

- (39) Weirich, K. L.; Banerjee, S.; Dasbiswas, K.; Witten, T. A.; Vaikuntanathan, S.; Gardel, M. L. Liquid behavior of cross-linked actin bundles. *Proc. Natl. Acad. Sci. U.S.A.* **2017**, *114*, 2131–2136.

- (40) Bonn, D.; Eggers, J.; Indekeu, J.; Meunier, J.; Rolley, E. Wetting and spreading. *Rev. Mod. Phys.* **2009**, *81*, 739–805.
- (41) Andrieu, C.; Beysens, D. A.; Nikolayev, V. S.; Pomeau, Y. Coalescence of sessile drops. *J. Fluid Mech.* **2002**, *453*, 427–438.
- (42) Cherry, B.; Holmes, C. Kinetics of wetting of surfaces by polymers. *J. Colloid Interface Sci.* **1969**, *29*, 174–176.
- (43) Van Oene, H.; Chang, Y. F.; Newman, S. The Rheology of Wetting By Polymer Melts. *J. Adhes.* **1969**, *1*, 54–68.
- (44) Varma, T.; Sharma, L.; Varma, S. Kinetics of spreading of a spherical droplet on a smooth surface. *Surf. Sci.* **1974**, *45*, 205–212.
- (45) de Ruijter, M. J.; De Coninck, J.; Blake, T. D.; Clarke, A.; Rankin, A. Contact Angle Relaxation during the Spreading of Partially Wetting Drops. *Langmuir* **1997**, *13*, 7293–7298.
- (46) Blake, T. D. The physics of moving wetting lines. *J. Colloid Interface Sci.* **2006**, *299*, 1–13.
- (47) Seveno, D.; Vaillant, A.; Rioboo, R.; Adão, H.; Conti, J.; De Coninck, J. Dynamics of Wetting Revisited. *Langmuir* **2009**, *25*, 13034–13044.
- (48) Blake, T. D.; Clarke, A.; De Coninck, J.; de Ruijter, M. J. Contact Angle Relaxation during Droplet Spreading: Comparison between Molecular Kinetic Theory and Molecular Dynamics. *Langmuir* **1997**, *13*, 2164–2166.
- (49) Blake, T. D.; de Coninck, J. The influence of solid-liquid interactions on dynamic wetting. *Adv. Colloid Interface Sci.* **2002**, *96*, 21–36.
- (50) Snoeijer, J. H.; Andreotti, B. Moving Contact Lines: Scales, Regimes, and Dynamical Transitions. *Annu. Rev. Fluid Mech.* **2013**, *45*, 269–292.
- (51) Voinov, O. V. Hydrodynamics of wetting. *Fluid Dyn.* **1977**, *11*, 714–721.
- (52) Xu, X.; Ma, L. Analysis of the effects of evaporative cooling on the evaporation of liquid droplets using a combined field approach. *Sci. Rep.* **2015**, *5*, No. 8614.
- (53) Dunn, G. J.; Wilson, S. K.; Duffy, B. R.; David, S.; Sefiane, K. The strong influence of substrate conductivity on droplet evaporation. *J. Fluid Mech.* **2009**, *623*, 329.
- (54) Lopes, M. C.; Bonaccorso, E.; Gambaryan-Roisman, T.; Stephan, P. Influence of the substrate thermal properties on sessile droplet evaporation: Effect of transient heat transport. *Colloids Surf., A* **2013**, *432*, 64–70.
- (55) Bazargan, V.; Stoeber, B. Effect of substrate conductivity on the evaporation of small sessile droplets. *Phys. Rev. E* **2016**, *94*, No. 033103.
- (56) Popov, Y. O. Evaporative deposition patterns: Spatial dimensions of the deposit. *Phys. Rev. E* **2005**, *71*, No. 036313.
- (57) Johnson, R. E.; Dettre, R. H. Contact Angle Hysteresis. III. Study of an Idealized Heterogeneous Surface. *J. Phys. Chem.* **1964**, *68*, 1744–1750.
- (58) Andersen, J. V.; Bréchet, Y. Pinning of a solid-liquid-vapor interface by stripes of obstacles. *Phys. Rev. E* **1996**, *53*, 5006–5010.
- (59) Brandon, S.; Marmur, A. Simulation of Contact Angle Hysteresis on Chemically Heterogeneous Surfaces. *J. Colloid Interface Sci.* **1996**, *183*, 351–355.
- (60) Schäffer, E.; Wong, P.-z. Dynamics of Contact Line Pinning in Capillary Rise and Fall. *Phys. Rev. Lett.* **1998**, *80*, 3069–3072.
- (61) Ondarçuhu, T.; Piednoir, A. Pinning of a Contact Line on Nanometric Steps during the Dewetting of a Terraced Substrate. *Nano Lett.* **2005**, *5*, 1744–1750.
- (62) Debuissou, D.; Merlen, A.; Senez, V.; Arscott, S. Stick–Jump (SJ) Evaporation of Strongly Pinned Nanoliter Volume Sessile Water Droplets on Quick Drying, Micropatterned Surfaces. *Langmuir* **2016**, *32*, 2679–2686.
- (63) de Coninck, J.; Fernández Toledano, J. C.; Dunlop, F.; Huillet, T. Pinning of a drop by a junction on an incline. *Phys. Rev. E* **2017**, *96*, No. 042804.